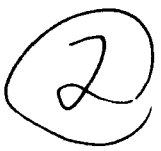

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Surface-Induced Static Undulations in Multilayer Films of Liquid-Crystalline Polymers

by

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13. ABSTRACT *(Maximum 200 words)*

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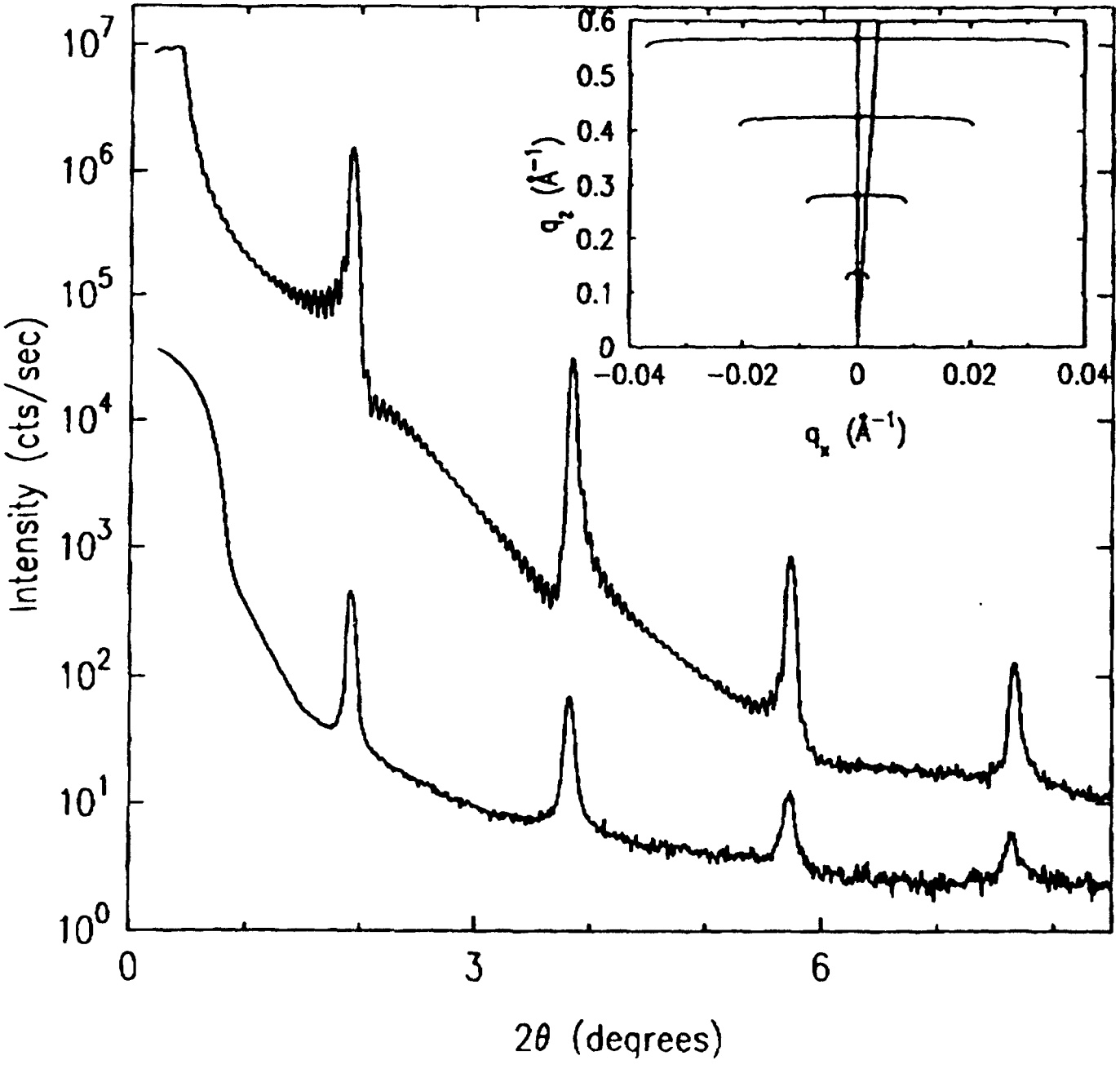
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q_z (Å^{-1})
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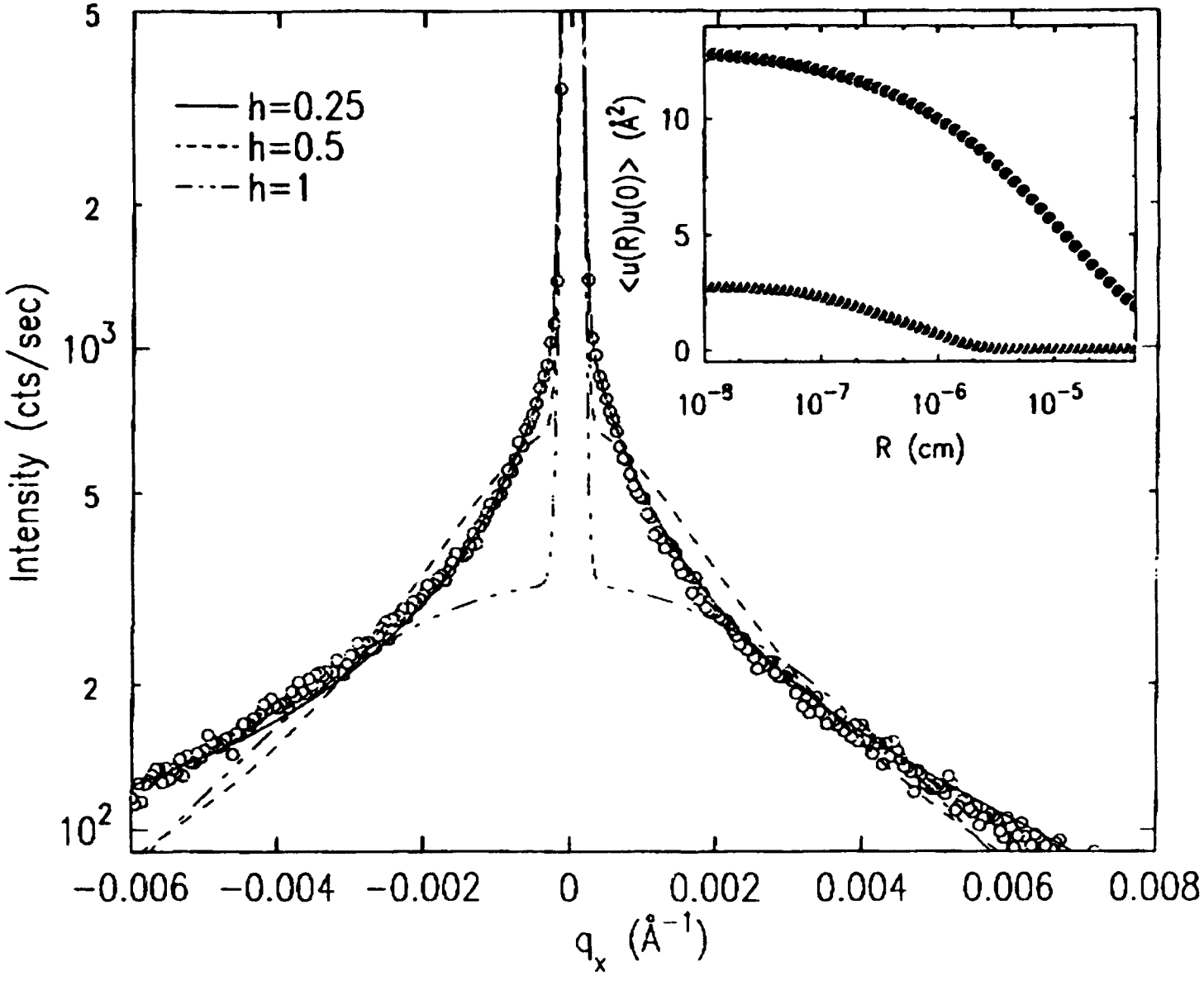
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10^2
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q_x (Å^{-1})
<u(R)u(0)> (Å^2)
10
5
0
10^{-8} 10^{-7} 10^{-6} 10^{-5}
R (cm)

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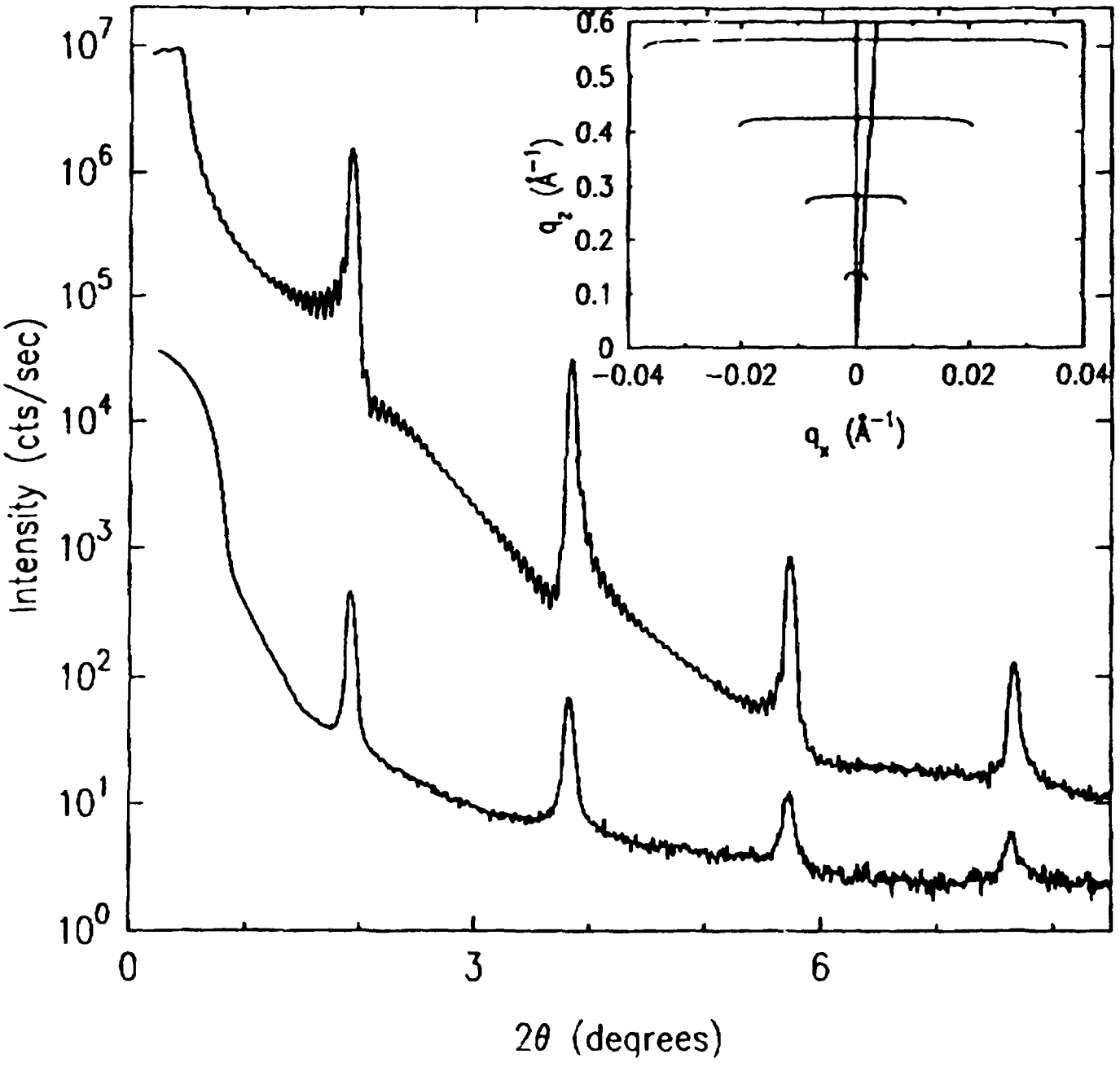
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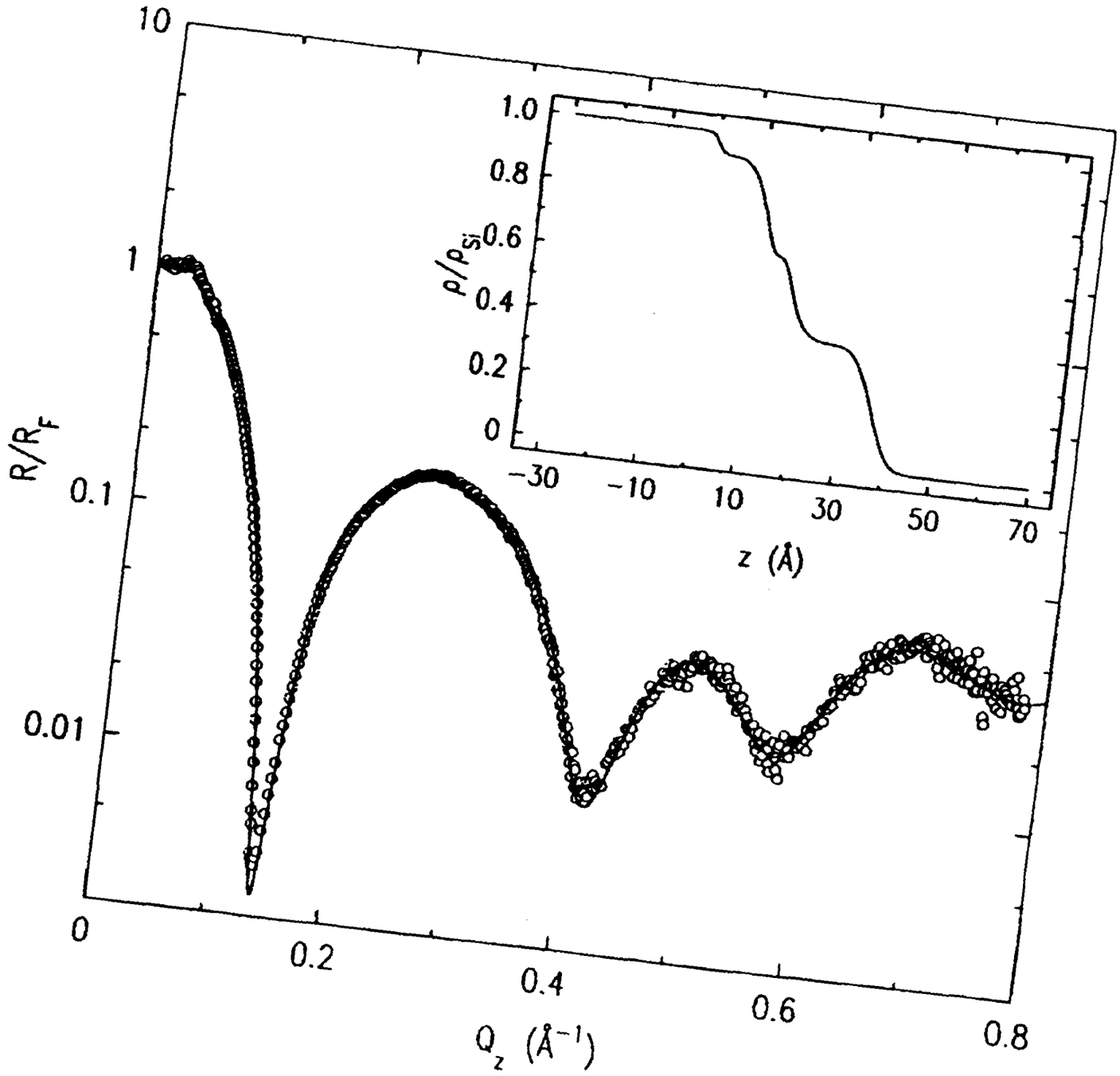
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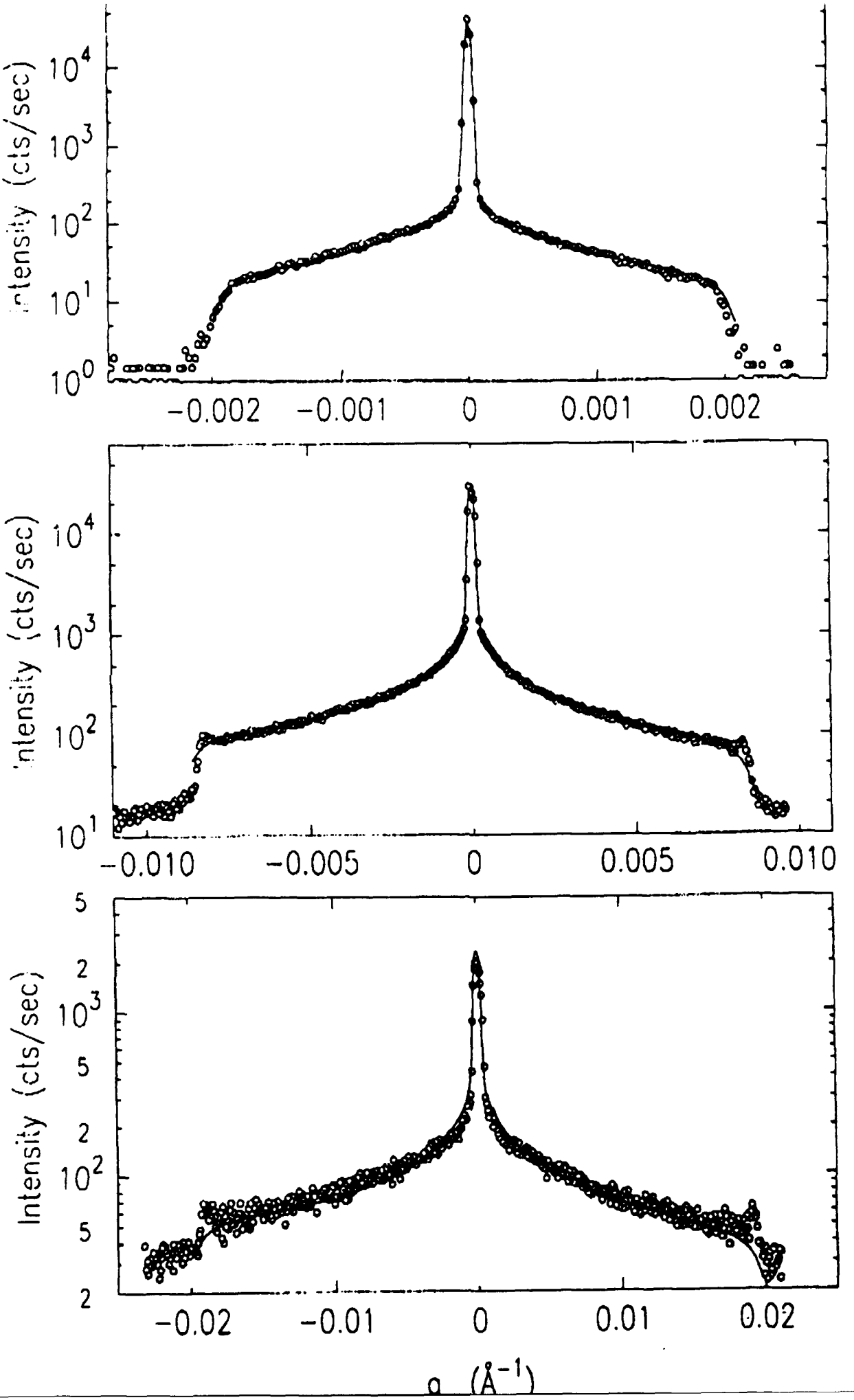
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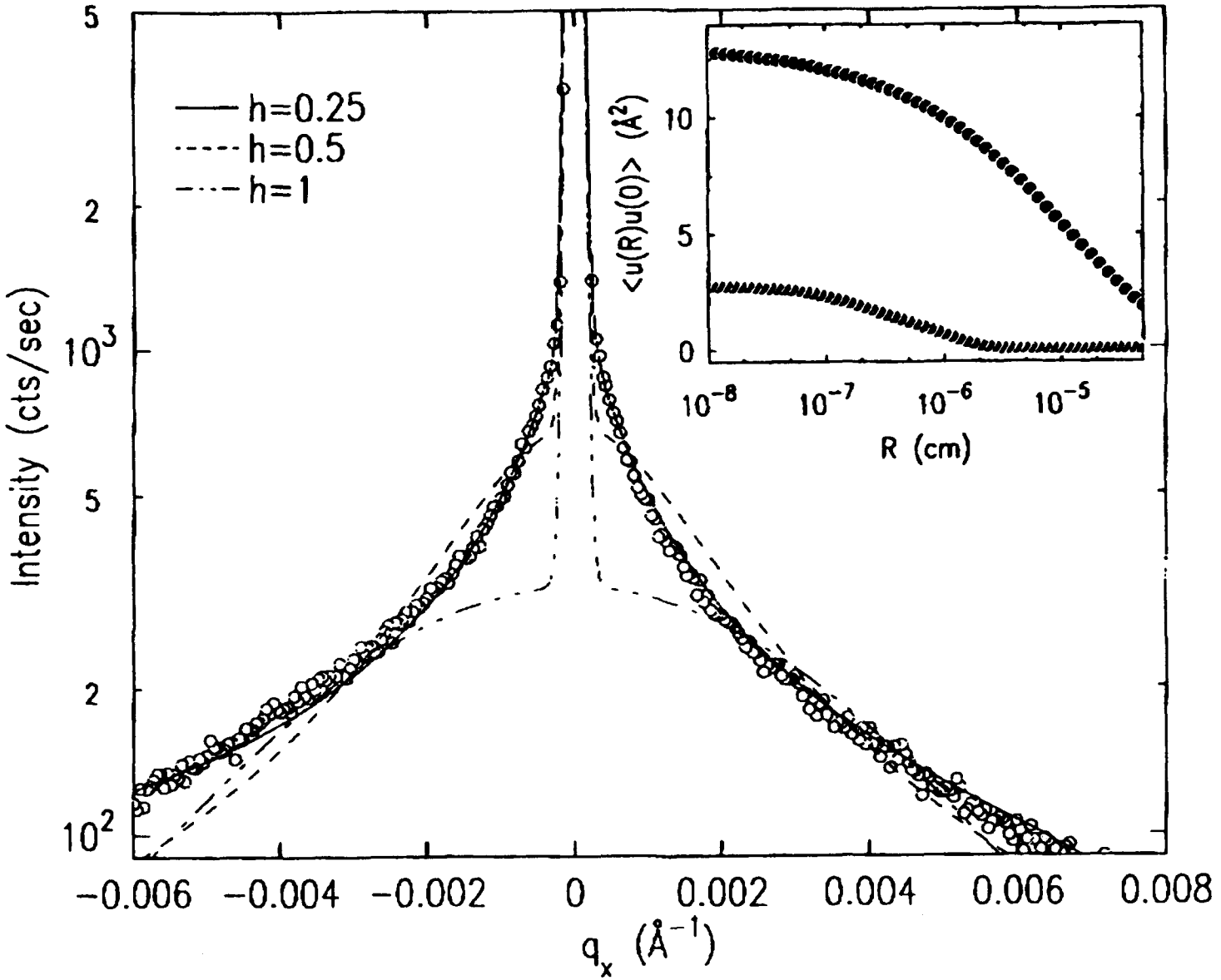
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0
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10^{-7}
10^{-6}
10^{-5}
R (cm)

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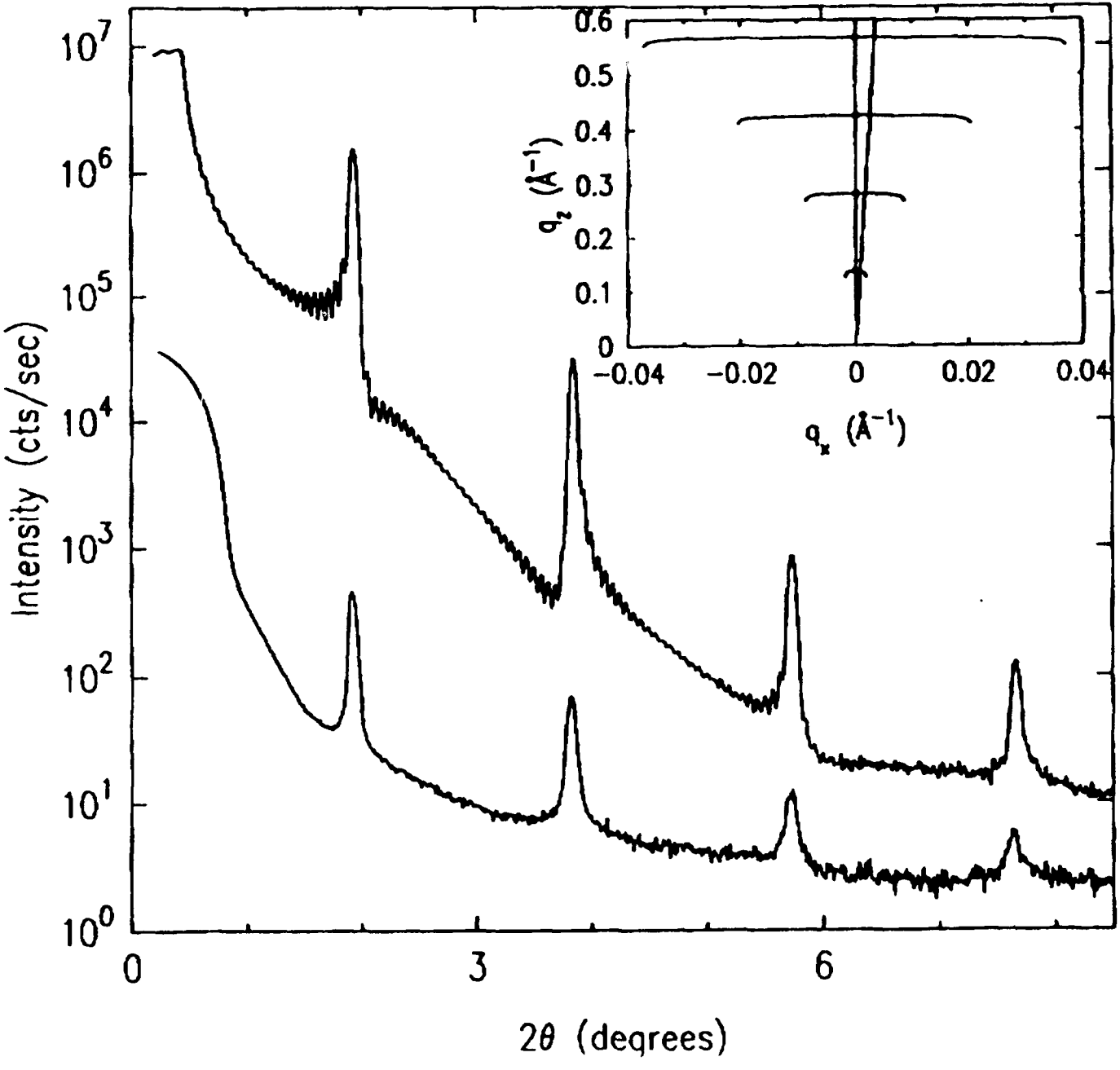
Intensity (cts/sec)
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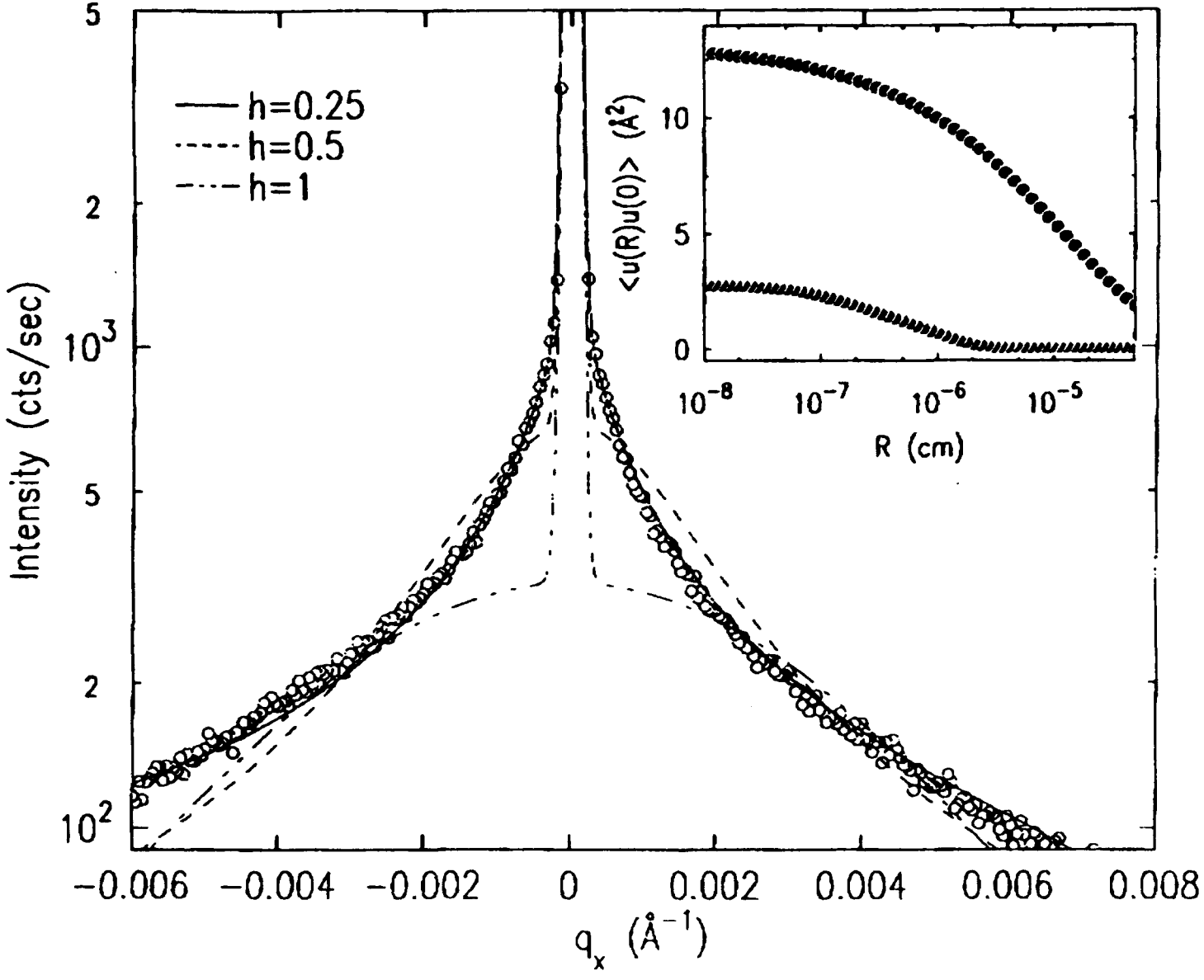
10^6
10^5
10^4
10^3
10^2
10^1
10^0
0
3
6
2θ (degrees)
0.6
0.5
0.4
0.3
0.2
0.1
0
q_z (Å^{-1})
−0.04
−0.02
0
0.02
0.04
q_x (Å^{-1})

Fig. 1


h=0.25
h=0.5
h=1
Intensity (cts/sec)
5
2
10^3
5
2
10^2
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q_x (Å^{-1})
<u(R)u(0)> (Å^2)
10
5
0
10^{-8} 10^{-7} 10^{-6} 10^{-5}
R (cm)

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Surface-Induced Static Undulations in Multilayer Films of Liquid-Crystalline Polymers

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Abstract

The first detailed study of surface-induced undulations in a liquid-crystalline polymer is presented. By examining the non-specular diffuse scattering from a 30-layer film of ferroelectric liquid-crystal polymer it is shown that the layer fluctuations are induced by the roughness of the film/substrate interface. This is in contrast to the case of free-standing films wherein thermal fluctuations play the *major* role.

Thin, smectic liquid-crystal films are of considerable current interest since they serve as model systems to study two-dimensional to three-dimensional crossover of inter- and intralayer order. For the most part, these studies have been on free-standing films. It is equally important to understand the influence of the interface between liquid-crystal films and the substrate. Film/vacuum and film/substrate interfaces are known to induce structural changes which are localized in the interfacial regions. On the other hand, such interfaces can also induce distinct thermodynamic phases as well as static undulations which penetrate into the interior of the film. The static undulations induced by roughness of the substrate surface has been studied in thin adsorbed films of cyclohexane. It was shown that for very thin films the substrate van der Waals interactions constrain the liquid surface to follow the static undulations of the substrate surface, while for thicker films the liquid surface structure is influenced mainly by thermally induced capillary waves. Early studies in homeotropically aligned smectic-

imperfections associated with the smectic layer order. Rocking scans through this diffuse scattering are shown in Fig. 2 for the first three Bragg peaks of Fig. 1. *These scans consist of broad diffuse scattering accompanied by a resolution limited peak due to specular reflection.* It may be recalled that Davidson and Levelut have reported similar diffuse bands perpendicular to the 00L axis for bulk samples of polysiloxane liquid-crystal polymers. These bands were attributed to undulations of longitudinally correlated rows of mesogens. The widths and amplitudes of the diffuse peaks of our LB multilayer shown in Fig. 1 are not consistent with such longitudinal undulations. We shall show that the diffuse scattering for the liquid-crystalline polymer is in fact due to static undulations of the smectic layers induced by the roughness of the substrate surface.

Quantitative analysis of the diffuse scattering of Fig. 2 is similar to that used by Sinha et al. for solid surfaces.[7] For a single rough surface, the scattered intensity is given by

$$S(\vec{q}) = \frac{1}{q_z^2} \iint_{S_o} dX dY e^{q_z^2 C(X,Y)} e^{-i(q_x X + q_y Y)} .$$

X and Y are the Cartesian separations of two points on the surface S_o with an average layer normal in the $\hat{z}$ direction. $C(X,Y)$ is the surface height-height correlation function. This is related to the average roughness across the sample $g(X,Y)=\langle[z(X,Y)-z(0)]^2\rangle$. For many isotropic solid surfaces $g(R=(X^2+Y^2)^{1/2}) = AR^{2h}$ describing so-called self-affine roughness[18]. $h = D_H-3$, where D_H is the fractal dimension of the surface. For systems of finite size (and measurement techniques with limited spatial resolution) $g(R) \rightarrow 2\sigma^2$ for large R, where σ is the rms roughness of the surface. A functional form satisfying these limits is

$$g(R) = 2\sigma^2 [1 - e^{-(R/\xi)^{2h}}].$$

ξ is a long-distance cutoff. Using this form for $g(R)$,

$$C(R) = \sigma^2 e^{-(R/\xi)^{2h}} .$$

$F(q_z)$ was determined by the specular portion of the rocking curves, the rms interlayer roughness σ, and the resolution function of the spectrometer. The fits are shown as solid lines in Fig. 2. *The agreement is excellent for all three rocking curves.* The data for all three quasi-Bragg peaks in Fig. 2 were fit using only *three significant adjustable parameters:* σ ξ and h.. Their values obtained from best fits are 3.6±0.12 Å, 1327±18 Å and 0.25±0.05, respectively. It should be mentioned that the wide range of q_z and q_x for the rocking curve data makes the fits extremely sensitive to the value of h. Figure 3, which plots the central portion of the 002 rocking curve with fits for three different values of h, demonstrates this sensitivity.

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